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(54) Title: LUNG REDUCTION DEVICE, SYSTEM, AND METHOD

(57) Abstract: A device, system, and method provides for lung size reduction by permanently collapsing at least a portion of a lung. The lung portion may be collapsed by obstructing the air passageway which communicates the lung portion to be collapsed. The air passageway may be obstructed by an obstructing member which precludes airflow in either direction or with a one-way valve which permits air to be exhaled from the lung portion while precluding air from being inhaled into the lung portion. In addition, a vacuum may be pulled within the lung portion to be collapsed for collapsing the lung portion and while the lung portion is collapsed the obstructing member may be placed in the air passageway to maintain the lung portion in a permanently collapsed state.

LUNG REDUCTION DEVICE, SYSTEM, AND METHOD

BACKGROUND OF THE INVENTION

5 The present invention is generally directed to a device, system, and method for treating Chronic Obstructive Pulmonary Disease (COPD). The present invention is more particularly directed to such a device, system and method which provide lung size reduction without requiring invasive surgery.

10 Chronic Obstructive Pulmonary Disease (COPD) has become a major cause of morbidity and mortality in the United States over the last three decades. COPD is characterized by the presence of airflow obstruction due to chronic bronchitis or emphysema. The airflow obstruction in COPD is due largely to structural abnormalities in the smaller airways. Important causes are inflammation, fibrosis, goblet cell metaplasia, and smooth muscle hypertrophy in terminal bronchioles.

15 The incidence, prevalence, and health-related costs of COPD are on the rise. Mortality due to COPD is also on the rise. In 1991 COPD was the fourth leading cause of death in the United States and had increased 33% since 1979.

20 COPD affects the patient's whole life. It has three main symptoms: cough; breathlessness; and wheeze. At first, breathlessness may be noticed when running for a bus, digging in the garden, or walking up hill. Later, it may be noticed when simply walking in the kitchen. Over time, it may occur with less and less effort until it is present all of the time.

25 COPD is a progressive disease and currently has no cure. Current treatments for COPD include the prevention of further respiratory damage, pharmacotherapy, and surgery. Each is discussed below.

 The prevention of further respiratory damage entails the adoption of a healthy lifestyle. Smoking cessation is believed to be the single most important therapeutic intervention. However, regular exercise and weight control are also important. Patients whose symptoms restrict their daily activities or who otherwise have an
30 impaired quality of life may require a pulmonary rehabilitation program including

ventilatory muscle training and breathing retraining. Long-term oxygen therapy may also become necessary.

Pharmacotherapy may include bronchodilator therapy to open up the airways as much as possible or inhaled β -agonists. For those patients who respond poorly to the foregoing or who have persistent symptoms, Ipratropium bromide may be indicated. Further, courses of steroids, such as corticosteroids, may be required. Lastly, antibiotics may be required to prevent infections and influenza and pneumococcal vaccines may be routinely administered. Unfortunately, there is no evidence that early, regular use of pharmacotherapy will alter the progression of COPD.

About 40 years ago, it was first postulated that the tethering force that tends to keep the intrathoracic airways open was lost in emphysema and that by surgically removing the most affected parts of the lungs, the force could be partially restored. Although the surgery was deemed promising, the procedure was abandoned.

The lung volume reduction surgery (LVRS) was later revived. In the early 1990's, hundreds of patients underwent the procedure. However, the procedure has fallen out of favor due to the fact that Medicare stopped reimbursing for LVRS. Unfortunately, data is relatively scarce and many factors conspire to make what data exists difficult to interpret. The procedure is currently under review in a controlled clinical trial. However, what data does exist tends to indicate that patients benefited from the procedure in terms of an increase in forced expiratory volume, a decrease in total lung capacity, and a significant improvement in lung function, dyspnea, and quality of life.

Improvements in pulmonary function after LVRS have been attributed to at least four possible mechanisms. These include enhanced elastic recoil, correction of ventilation/perfusion mismatch, improved efficiency of respiratory musculature, and improved right ventricular filling.

Lastly, lung transplantation is also an option. Today, COPD is the most common diagnosis for which lung transplantation is considered. Unfortunately, this consideration is given for only those with advanced COPD. Given the limited availability of donor organs, lung transplant is far from being available to all patients.

In view of the foregoing, there is a need in the art for a new and improved therapy for COPD. More specifically, there is a need for such a therapy which provides more permanent results than pharmacotherapy while being less invasive and traumatic than LVRS. The present invention is directed to a device, system, and
5 method which provide such an improved therapy for COPD.

SUMMARY OF THE INVENTION

The present invention provides a method of reducing the size of a lung including the step of permanently collapsing at least a portion of the lung. In
10 accordance with a first embodiment, the lung may be collapsed by obstructing an air passageway communicating with the lung portion to be collapsed. The air passageway may be obstructed by placing an obstructing member in the air passageway. The obstructing member may be a plug-like device which precludes air flow in both directions or a one-way valve which permits air to be exhaled from
15 the lung portion to be collapsed while precluding air from being inhaled into the lung portion. Once the air passageway is sealed, the residual air within the lung will be absorbed over time to cause the lung portion to collapse.

In accordance with a further embodiment of the present invention, the lung portion may be collapsed by inserting a conduit into the air passageway
20 communicating with the lung portion to be collapsed, pulling a vacuum in the lung portion through the conduit to collapse the lung portion, and maintaining the lung portion in a collapsed state. The lung portion may be maintained in a collapsed state by sealing the air passageway with an obstructing member or by placing a one-way valve in the air passageway. To efficiently pull the vacuum in the lung portion to
25 be collapsed, the space between the outer surface of the conduit and the inner surface of the air passageway may be sealed as the vacuum is pulled. Preferably, the air passageway is sealed while the lung portion is collapsed.

The present invention further provides a device for reducing the size of a lung. The device includes an obstructing member insertable into an air passageway
30 communicating with a portion of the lung to be reduced in size and having an inner dimension. The obstructing member has an outer dimension for continuous contact

with the air passageway inner dimension and sealing the air passageway upon placement in the air passageway for collapsing the portion of the lung and reducing the size of the lung. The obstructing member may be formed of resilient material so as to be collapsible for initial insertion into the air passageway in a collapsed

5 condition and releasable to define the outer dimension upon placement in the air passageway. In accordance with a further embodiment of the present invention, the obstructing member may include a one-way valve to permit exhaled air to flow from the lung portion while precluding inhaled air from flowing into the lung portion.

The present invention further provides a system for reducing the size of a
10 lung. This system includes a conduit configured to be passed down a trachea, into a bronchus communicating with the trachea, and into an air passageway communicating the bronchus with a lung portion to be reduced in size. The system further includes an obstructing member configured to be guided through the conduit into the air passageway for placement in the air passageway and sealing the air
15 passageway for collapsing the lung portion. The conduit preferably has an outer dimension smaller than the inner dimension of the passageway and a sealing member seals the space between the conduit outer dimension and the air passageway inner dimension as the vacuum is pulled the system may further include a vacuum source for pulling a vacuum in the lung portion through the conduit
20 prior to placement of the obstructing member. The sealing member may be an inflatable member. The conduit may include a first channel pulling the vacuum and for guiding the obstructing member into position and a second channel for inflating the inflatable sealing member.

25

BRIEF DESCRIPTION OF THE DRAWINGS

The features of the present invention which are believed to be novel are set forth with particularity in the appended claims. The invention, together with further objects and advantages thereof, may best be understood by making reference to the following description taken in conjunction with the accompanying drawings, in the
30 several figures of which like referenced numerals identify identical elements, and wherein:

Figure 1 is a simplified sectional view of a thorax illustrating a healthy respiratory system;

Figure 2 is a sectional view similar to FIG. 1 but illustrating a respiratory system suffering from COPD and the execution of a first step in treating the COPD condition by reducing the size of a lung portion in accordance with the present invention;

Figure 3 is a perspective view, partly in section, and to an enlarged scale, illustrating an intermediate step in the treatment;

Figure 4 is a partial perspective view of a conduit which may be utilized in practicing the present invention;

Figure 5 is a perspective view, to an enlarged scale, illustrating the guiding of an obstructing member through the conduit for sealing a lung portion in accordance with the present invention;

Figure 6 is a partial exploded view of FIG. 5;

Figure 7 is a perspective view, partly in section, and to an enlarged scale, illustrating an obstructing member positioned in an air passageway for sealing the lung portion;

Figure 8 is a perspective view, to an enlarged scale, of an obstructing member configured in accordance with the present invention;

Figure 9 is a perspective view, to an enlarged scale, of the obstructing member of FIG. 8 having a reinforcing rib;

Figure 10 is a perspective view, to an enlarged scale, illustrating the obstructing member of FIG. 6 in a collapsed condition in accordance with further aspects of the present invention;

Figure 11 is a perspective view, to an enlarged scale, of another obstructing member embodying the present invention in the form of a one-way valve;

Figure 12 is a sectional view, to an enlarged scale, of another one-way valve obstructing member embodying the present invention in a closed position; and

Figure 13 is a sectional view, to an enlarged scale, of the one-way valve of FIG. 12 illustrated in an open condition.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

Referring now to FIG. 1, it is a sectional view of a healthy respiratory system. The respiratory system 20 resides within the thorax 22 which occupies a space defined by the chest wall 24 and the diaphragm 26.

5 The respiratory system 20 includes the trachea 28, the left mainstem bronchus 30, the right mainstem bronchus 32, the bronchial branches 34, 36, 38, 40, and 42 and sub-branches 44, 46, 48, and 50. The respiratory system 20 further includes left lung lobes 52 and 54 and right lung lobes 56, 58, and 60. Each bronchial branch and sub-branch communicates with a respective different portion of
10 a lung lobe, either the entire lung lobe or a portion thereof. As used herein, the term "air passageway" is meant to denote either a bronchial branch or sub-branch which communicates with a corresponding individual lung lobe or lung lobe portion to provide inhaled air thereto or conduct exhaled air therefrom.

 Characteristic of a healthy respiratory system is the arched or inwardly
15 arcuate diaphragm 26. As the individual inhales, the diaphragm 26 straightens to increase the volume of the thorax 22. This causes a negative pressure within the thorax. The negative pressure within the thorax in turn causes the lung lobes to fill with air. When the individual exhales, the diaphragm returns to its original arched condition to decrease the volume of the thorax. The decreased volume of the thorax
20 causes a positive pressure within the thorax which in turn causes exhalation of the lung lobes.

 In contrast to the healthy respiratory system of FIG. 1, FIG. 2 illustrates a respiratory system suffering from COPD. Here it may be seen that the lung lobes 52, 54, 56, 58, and 60 are enlarged and that the diaphragm 26 is not arched but
25 substantially straight. Hence, this individual is incapable of breathing normally by moving the diaphragm 28. Instead, in order to create the negative pressure in the thorax 22 required for breathing, this individual must move the chest wall outwardly to increase the volume of the thorax. This results in inefficient breathing causing these individuals to breathe rapidly with shallow breaths.

30 It has been found that the apex portion 62 and 66 of the upper lung lobes 52 and 56, respectively, are most affected by COPD. Hence, the preferred embodiment

will be described for treating the apex 66 of the right, upper lung lobe 56. However, as will be appreciated by those skilled in the art, the present invention may be applied to any lung portion without departing from the present invention.

The device, system, and method of the present invention treats COPD by
5 deriving the benefits of lung volume reduction surgery without the need of performing lung volume reduction surgery. As will be seen hereinafter, the present invention contemplates permanent collapse of a lung portion or lung portions most affected. This leaves extra volume within the thorax for the diaphragm to assume its arched state for acting upon the remaining healthier lung tissue. As previously
10 mentioned, this should result in improved pulmonary function due to enhanced elastic recoil, correction of ventilation/perfusion mismatch, improved efficiency of respiratory musculature, and improved right ventricle filling.

In accordance with this embodiment of the present invention, the COPD treatment is initiated by feeding a conduit or catheter 70 down the trachea 28, into
15 the right mainstem bronchus 32, into the bronchial branch 42 and into and terminating within the sub-branch 50. The sub-branch 50 is the air passageway which communicates with the lung portion 66 to be treated.

The catheter 70 is preferably formed of flexible material such as polyethylene. Also, the catheter 70 is preferably preformed with a bend 72 to assist the feeding of
20 the catheter from the right mainstem bronchus 32 into the bronchial branch 42.

Referring now to FIG. 3, here it may be seen that the catheter includes an inflatable sealing member 74. The inflatable sealing member 74 is inflated within the sub-branch 50 to seal the space in-between the inner dimension 51 of the passageway 50 and the outer dimension 75 of the catheter 70. For inflating the
25 inflatable member 74 and pulling a vacuum within lung portion 66, the catheter 70 is coupled to a pump 80. As may be seen in FIG. 4, the catheter 70 includes a main channel 82 through which the vacuum in lung portion 66 is pulled and a minor channel 84 which is utilized for inflating the inflatable member 74.

To establish the vacuum in lung portion 66, the inflatable member 74 is first
30 inflated. Thereafter, the vacuum is pulled through the main channel 82 of the catheter 70 to pull the vacuum in lung portion 66.

Referring now to FIG. 5, here it may be seen that the lung portion 66, due to the vacuum pulled by the pump 80 and catheter 70 of FIG. 3, has collapsed from its initial state indicated by the dashed line 86 to the solid line 88. With the lung portion 66 thus collapsed, and while the lung portion 66 is collapsed, an obstructing member 90 is guided through the main channel of the conduit 70 by a stylet wire 92. This may be seen in greater detail in FIG. 6. As will be seen hereinafter with specific reference to FIGS. 8-13, the obstructing or sealing member 90 is formed of resilient or collapsible material to enable the obstructing member 90 to be fed through the conduit 70 in a collapsed state. The stylet 92 is used to push the obstructing member 90 to the end 77 of the catheter 70 for placing the obstructing member 90 within the air passageway 50 adjacent to the lung portion 66 to be permanently collapsed.

FIG. 7 illustrates the obstructing or sealing member in place within the air passageway 50. The sealing member 90 has expanded upon placement in the air passageway 50 to seal the air passageway 50. This causes the lung portion 66 to be maintained in a permanently collapsed state.

More specifically, the obstructing member 90 has an outer dimension 91 when expanded to enable continuous contact with the air passageway inner dimension 51. This seals the air passageway upon placement of the obstructing member 90 in the air passageway 50 for maintaining the lung portion 66 in the collapsed state.

Alternatively, the vacuum need not be pulled in the lung portion 66. Rather, the lung portion 66, in accordance with a further embodiment of the present invention, may be collapsed by sealing the air passageway 50 with the obstructing member 90. Over time, the air within the lung portion 66 will be absorbed by the body to result in the collapse of lung portion 66. In accordance with this embodiment of the present invention, the obstructing member 90 may be placed in the air passageway 50 by utilizing catheter 70 as previously described but without pulling a vacuum or employing the inflatable sealing member 74.

Referring now to FIG. 8, it illustrates the ceiling or obstructing member 90 in greater detail. The obstructing member 90 has a hollow cylindrical configuration.

More specifically, the obstructing member 90 includes a generally circular base 94 having an outer periphery 95. The obstructing member 90 further includes a circumferential generally cylindrical sidewall 96 which extends from the outer periphery 95 of the base 94. The generally cylindrical sidewall 96 has an outer surface 98 which defines the outer periphery 91 of the obstructing member as illustrated in FIG. 7.

As previously mentioned, the obstructing member 90 is formed of resilient material. For example, the obstructing member 90 may be formed from silicone rubber. This renders the obstructing member 90 collapsible as illustrated in FIGS. 5 and 6 for feeding the obstructing member 90 through the catheter 70 and causing the obstructing member 90 to expand when placed in the air passageway 50 for sealing the air passageway.

FIG. 9 illustrates the obstructing member 90 as described with respect to FIG. 8 but, in addition, includes an inner resilient reinforcement rib 100. The reinforcement rib 100 has a generally serpentine configuration and is in contact with the inner generally cylindrical surface 102 of the obstructing member 90. The reinforcement rib 100 serves to add increased structural integrity to the obstructing member 90. In addition, the base 94, the generally cylindrical sidewall 96, and the resilient reinforcement rib 100 are all collapsible. The reinforcement rib 100, when the obstructing member 90 is positioned in place within the air passageway 50, forces the cylindrical sidewall 96 radially outwardly to form the generally circular base 94 and the circumferential generally cylindrical sidewall 96.

FIG. 10 illustrates the obstructing member 90 in its fully collapsed state. Here it may be seen that the base 94, when collapsed, is generally conically shaped to assist in the placement of the collapsed obstructing member in the main channel of the conduit. Also, as can be clearly seen in FIG. 10, the resilient reinforcing rib 100 is collapsed and ready to expand the obstructing member 90 by forcing the cylindrical sidewall 96 radially outwardly.

Referring now to FIG. 11, it shows, in accordance with a further aspect of the present invention, an obstructing member 110 which is similar to the previously described obstructing member 90 but which includes a one-way valve. More

specifically, the obstructing member 110 includes the generally circular base 94, the generally cylindrical sidewall 96, and the reinforcement rib 100. In addition, the base 94 is slit as illustrated at 104 to form a valve structure. A tether 106 is connected between the reinforcement rib 100 and the base 94. Hence, when positioned in the air passageway 50, the one-way valve structure of the obstructing member 110 permits air to flow in the direction indicated by the arrow 108 but precludes airflow in the opposite direction. Hence, when the obstructing member 110 is placed in the air passageway 50, it is so placed in accordance with this aspect of the present invention that it permits air to be exhaled from the lung portion to be collapsed but precludes air from being inhaled into the lung portion to be collapsed.

The one-way valve obstructing member 110 may be utilized in accordance with the embodiment wherein a vacuum is pulled within the lung portion to be collapsed or in accordance with the alternative embodiment wherein the air passageway is obstructed without the pulling of a vacuum in the lung portion to be collapsed. Because the one-way valve obstructing member 110 permits exhaled air to flow therethrough, the lung portion to be collapsed will be collapsed more quickly due to the reduction in residual air within the lung portion to be collapsed.

Referring now to FIGS. 12 and 13, they illustrate another one-way valve obstructing member 120 which may be utilized in accordance with the present invention. Like the previous obstructing members, the one-way valve obstructing member 120 includes a generally circular base 94 and a circumferential generally cylindrical sidewall 96. The obstructing member 120 further includes the reinforcement rib 100. To form the valve, the base 94 includes a slit 122. On either side of the slit 122 is a tether 124 and 126 which extend to the resilient reinforcement rib 100. As a result, the one-way valve structure opens to air flow in the direction indicated by arrow 128 but precludes air flow in the opposite direction. The one-way valve obstructing member 120 may thus be employed in the same manner as the one-way valve 110 as described with reference to FIG. 11.

As can thus be seen from the foregoing, the present invention provides a device, system, and method for treating COPD by lung volume reduction. The lung volume reduction is achieved through the permanent collapsing of one or more lung

portions, or lobes, or portions of lobes. The foregoing is achieved without surgery.

Following the treatment, the lung tissue within the thorax will occupy a lesser volume than previously occupied providing room for the diaphragm to assume its arcuate state to assist in normal breathing and to achieve the benefits of lung volume

5 reduction.

While particular embodiments of the present invention have been shown and described, modifications may be made, and it is therefore intended in the appended claims to cover all such changes and modifications which fall within the true spirit and scope of the invention.

10

What is claimed is:

1. A method of reducing the size of a lung, the method including permanently collapsing at least a portion of the lung.

5

2. The method of claim 1 wherein the permanently collapsing step includes obstructing an air passageway communicating with the lung portion to be collapsed.

10 3. The method of claim 2 wherein the obstructing step includes placing an obstructing member in the air passageway.

4. The method of claim 2 wherein the obstructing step includes placing a one-way valve in the air passageway, the one-way valve permitting air to be exhaled
15 from the lung portion while precluding air from being inhaled into the lung portion.

5. The method of claim 1 wherein the permanently collapsing step includes inserting a conduit into the air passageway communicating with the lung portion to be collapsed, pulling a vacuum in the lung portion through the conduit to
20 collapse the lung portion, and maintaining the lung portion in a collapsed state.

6. The method of claim 5 wherein the conduit includes an outer surface, wherein the air passageway has an inner surface, and wherein the pulling a vacuum step includes sealing between the outer surface of the conduit and the inner surface
25 of the air passageway.

7. The method of claim 5 wherein the maintaining step includes sealing the lung portion from the air passageway while the lung portion is collapsed.

30 8. The method of claim 7 wherein the sealing step includes placing an obstructing member in the air passageway.

9. The method of claim 7 wherein the sealing step includes placing a one-way valve in the air-passageway, the one-way valve permitting air to be exhaled from the lung portion while precluding air from being inhaled into the lung portion.

5

10. A device for reducing the size of a lung comprising an obstructing member insertable into an air passageway communicating with a portion of the lung to be reduced in size and having an inner dimension, the obstructing member having an outer dimension for continuous contact with the air passageway inner dimension and sealing the air passageway upon placement in the air passageway for collapsing the portion of the lung and reducing the size of the lung.

10

11. The device of claim 10 wherein the obstructing member is formed of resilient material so as to be collapsible for initial insertion into the air passageway in a collapsed condition and releasable to define the outer dimension upon placement in the air passageway.

15

12. The device of claim 10 wherein the obstructing member includes a generally circular base having an outer periphery and a circumferential generally cylindrical sidewall extending from the outer periphery of the base, the generally cylindrical sidewall having an outer surface defining the outer dimension.

20

13. The device of claim 12 wherein the obstructing member is formed of resilient material.

25

14. The device of claim 13 wherein the obstructing member is formed of silicone rubber.

15. The device of claim 13 wherein the obstructing member sidewall includes an inner generally cylindrical surface and wherein the obstructing member

30

includes a resilient reinforcement rib in contact with the inner generally cylindrical surface.

5 16. The device of claim 15 wherein the reinforcement rib has a serpentine configuration.

10 17. The device if claim 15 wherein the obstructing member further includes a one-way valve within the generally circular base to permit exhaled air to flow from the lung portion while precluding inhaled air from flowing into the lung portion.

15 18. The device of claim 15 wherein the base, the generally cylindrical sidewall, and the resilient reinforcement rib of the obstructing member are all collapsible and wherein the resilient reinforcement rib forces the cylindrical sidewall radially outwardly when the obstructing member is positioned in the air passageway to form the generally circular base and circumferential generally cylindrical sidewall.

20 19. The device of claim 10 wherein the obstructing member is a one-way valve to permit exhaled air to flow from the lung portion while precluding inhaled air from flowing into the lung portion.

25 20. A system for reducing the size of a lung comprising:
a conduit configured to be passed down a trachea, into a bronchus communicating with the trachea, and into an air passageway communicating the bronchus with a lung portion to be reduced in size; and
an obstructing member configured to be guided through the conduit into the air passageway for placement in the air passageway and sealing the air passageway for collapsing the lung portion.

30 21. The system of claim 20 further including a vacuum source for pulling a vacuum in the lung portion prior to placement of the obstructing member.

22. The system of claim 21 wherein the vacuum source is adapted to be coupled to the conduit for pulling the vacuum through the conduit.

5 23. The system of claim 22 wherein the air passageway has an inner dimension, wherein the conduit has an outer dimension smaller than the passageway inner dimension, and wherein the system further includes sealing means for sealing between the conduit outer dimension and the air passageway inner dimension.

10 24. The system of claim 23 wherein the sealing means comprises an inflatable member.

15 25. The system of claim 24 wherein the conduit includes a first channel for guiding the obstructing member and a second channel for inflating the inflatable member.

20 26. The system of claim 20 wherein the obstructing member is formed of resilient material so as to be collapsible for initial insertion through the conduit into the air passageway in a collapsed condition and releasably expandable to seal the air passageway upon placement in the air passageway.

25 27. The system of claim 20 wherein the obstructing member includes a generally circular base having an outer periphery and a circumferential generally cylindrical sidewall extending from the outer periphery of the base, the generally cylindrical sidewall having an outer surface for continuous engagement with the air passageway.

30 28. The system of claim 27 wherein the obstructing member is formed of resilient material.

29. The system of claim 28 wherein the obstructing member is formed of silicone rubber.

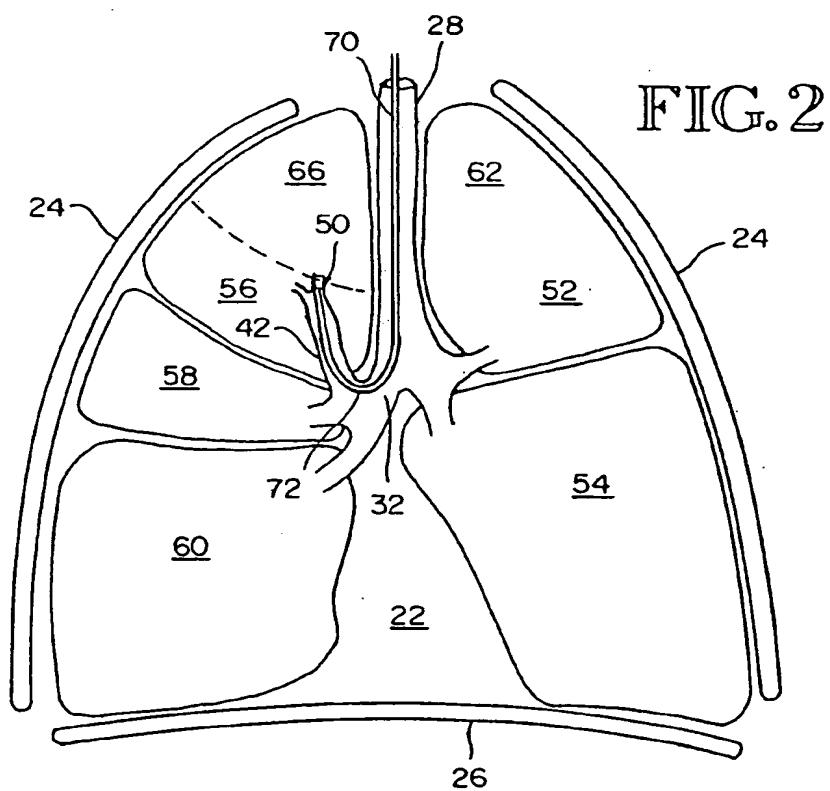
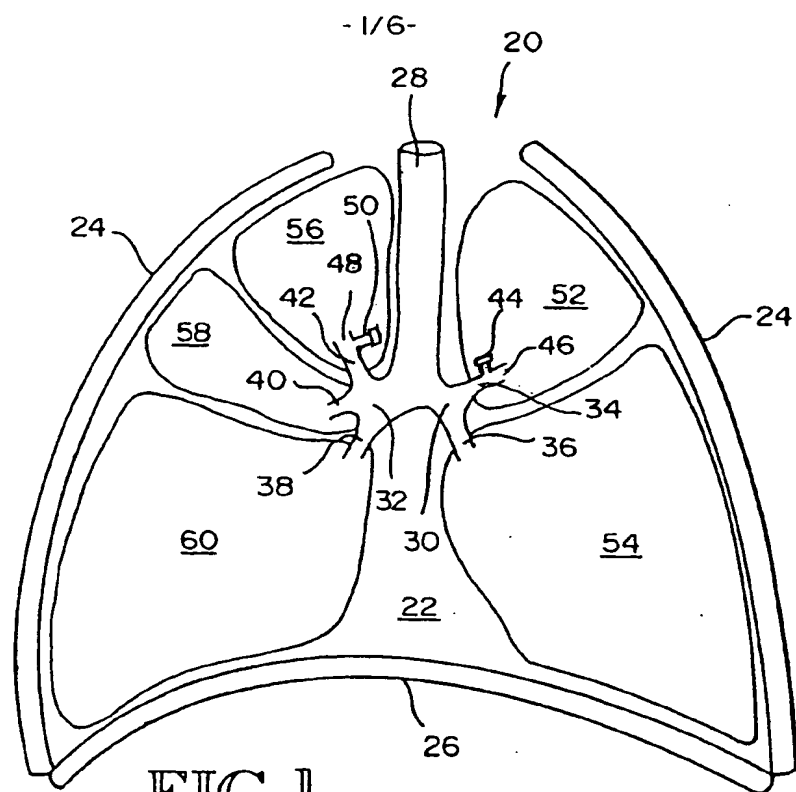
30. The system of claim 28 wherein the obstructing member sidewall
5 includes an inner generally cylindrical surface and wherein the obstructing member includes a resilient reinforcement rib in contact with the inner generally cylindrical surface.

31. The system of claim 30 wherein the reinforcement rib has a serpentine
10 configuration.

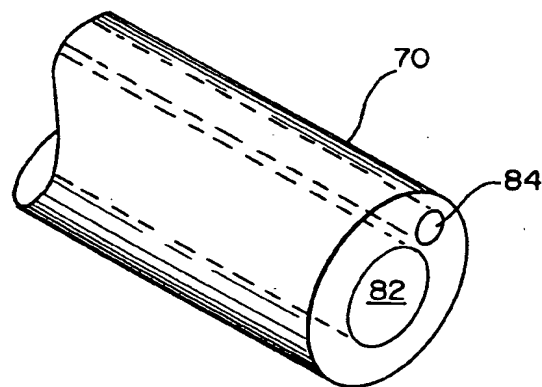
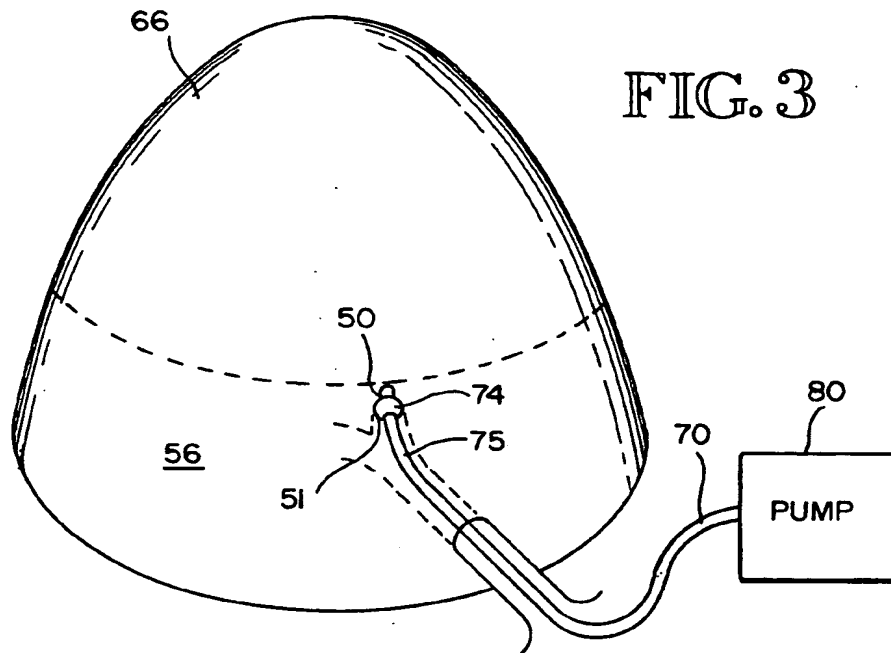
32. The system of claim 30 wherein the obstructing member further
includes a one-way valve within the generally circular base to permit exhaled air to
flow from the lung portion while precluding inhaled air from flowing into the lung
15 portion.

33. The system of claim 30 wherein the base, the generally cylindrical
sidewall, and the resilient reinforcement rib of the obstructing member are all
collapsible and wherein the resilient reinforcement rib forces the cylindrical sidewall
20 radially outwardly when the obstructing member is positioned in the air passageway
to form the generally circular base and circumferential cylindrical sidewall.

34. The system of claim 20 wherein the obstructing member is a one-way
valve to permit exhaled air to flow from the lung portion while precluding inhaled air
25 from flowing into the lung portion.



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-3/6-

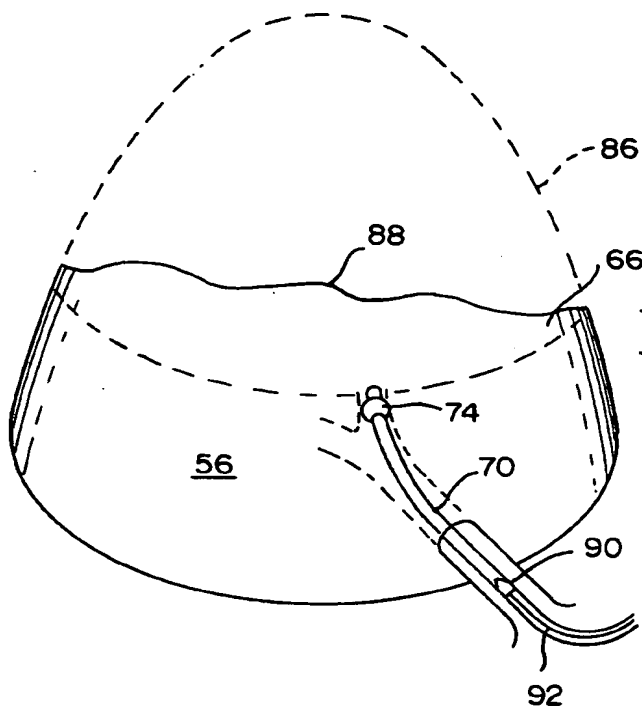


FIG. 5

FIG. 6

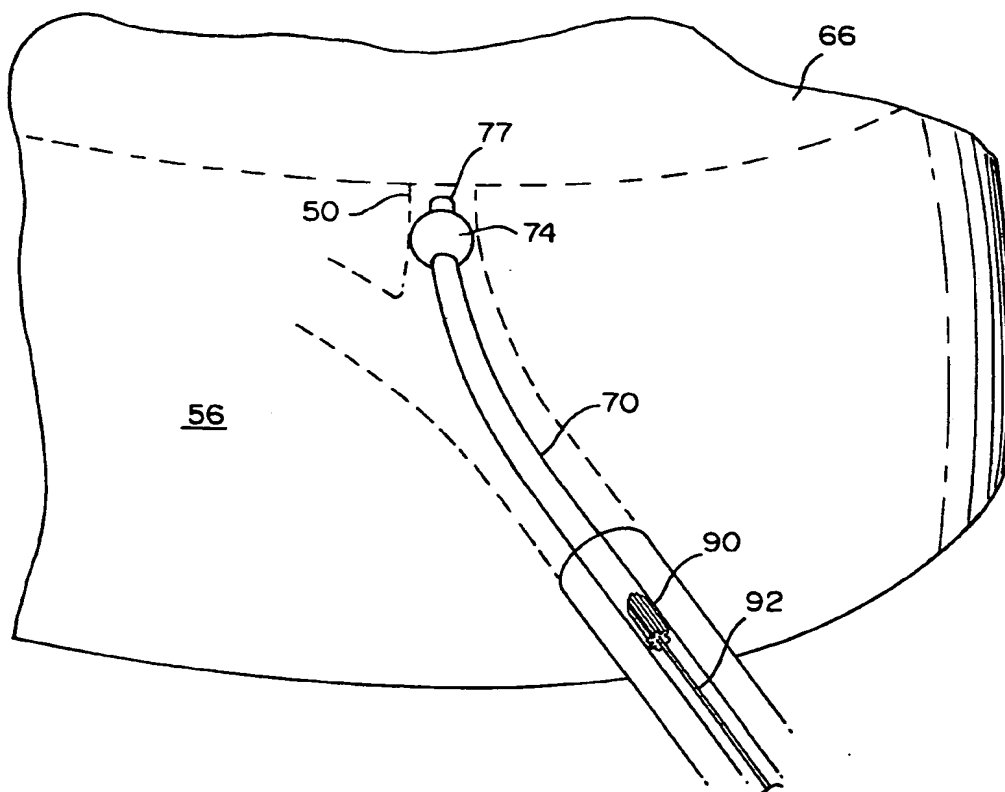


FIG. 7

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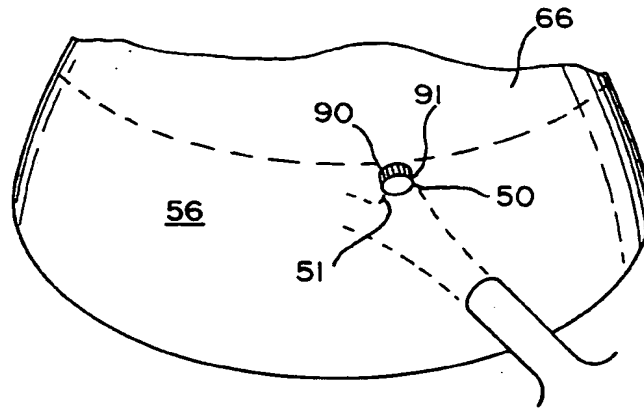


FIG. 8

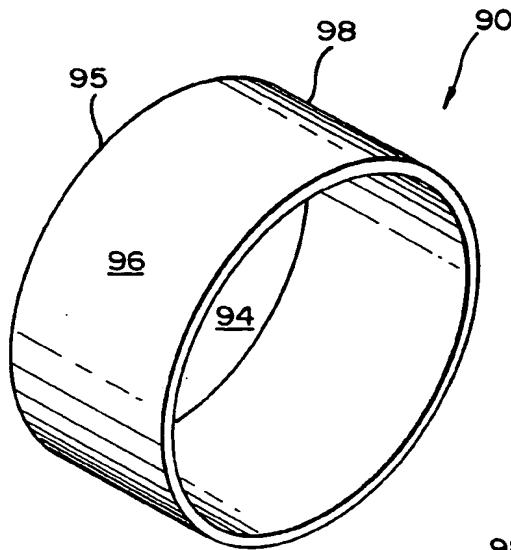
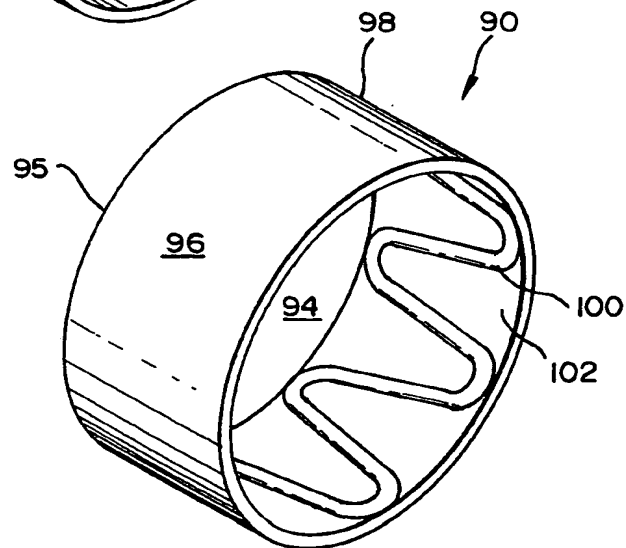
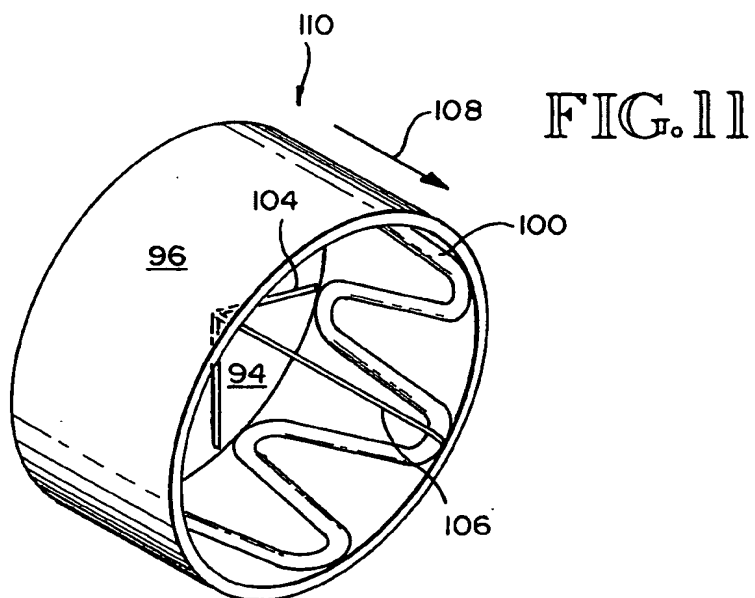
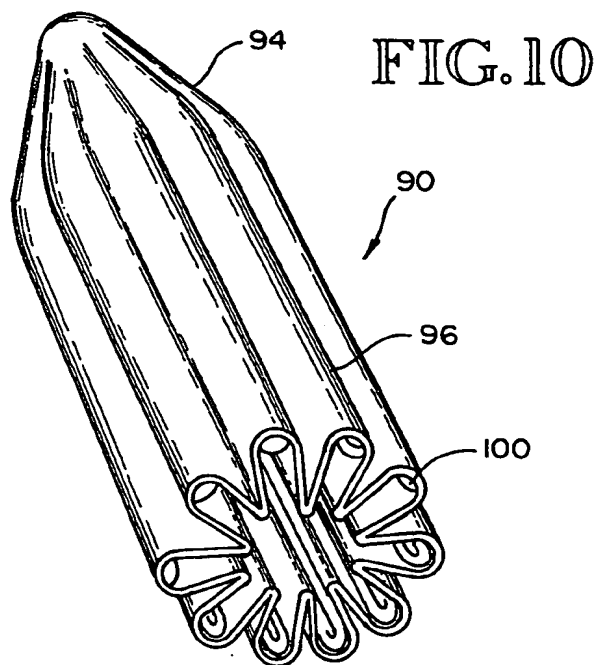


FIG. 9



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-6/6-

FIG. 12

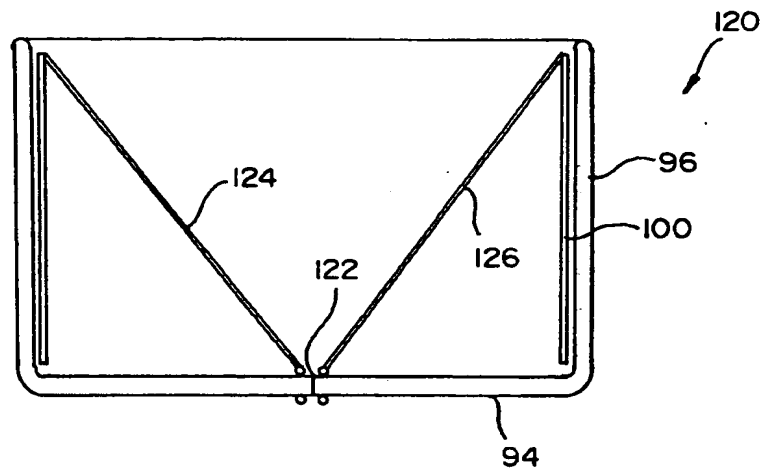
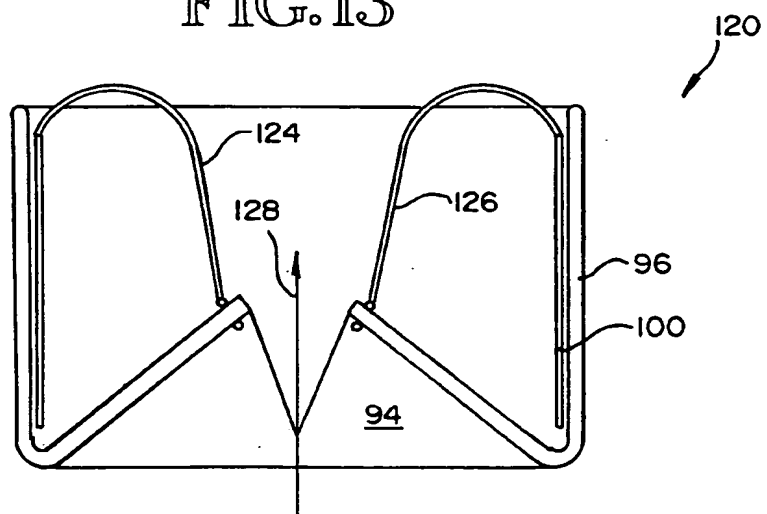


FIG. 13



INTERNATIONAL SEARCH REPORT

International application No.
PCT/US00/40701

A. CLASSIFICATION OF SUBJECT MATTER IPC(7) : A61F 11/00 US CL : 606/108; 128/898 According to International Patent Classification (IPC) or to both national classification and IPC																										
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) U.S. : 606/108, 200, 213, 217; 604/246, 247, 249, 256, 48; 623/1.26, 1.24, 23.70; 128/898 Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) EASR search terms: chronic obstructive pulmonary disease, collaps, lung, obstruct, air, reduced, size, copd																										
C. DOCUMENTS CONSIDERED TO BE RELEVANT																										
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.																								
X	US 5,306,234, A (JOHNSON) 26 April 1994, col. 2, lines 60-65	1																								
X	US 3,617,060 A (IEZZI) 2 November 1971, col. 1, lines 1-8.	1																								
X --- Y	US 5,855,601 A (BESSLER et al) 5 January 1999, see complete document.	10-13, 15-20, 26-28, 30-34 --- 14, 29																								
X, P	US 6,027,525 A (SUH et al) 22 February 2000, see complete document.	10-13, 15-20, 26-28, 30-34																								
<input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C. <input type="checkbox"/> See patent family annex.																										
<table border="0"> <tr> <td>* Special categories of cited documents:</td> <td>*T</td> <td>later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</td> </tr> <tr> <td>*A</td> <td>document defining the general state of the art which is not considered to be of particular relevance</td> <td></td> </tr> <tr> <td>*E</td> <td>earlier document published on or after the international filing date</td> <td>*X</td> </tr> <tr> <td>*L</td> <td>document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</td> <td>document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</td> </tr> <tr> <td>*O</td> <td>document referring to an oral disclosure, use, exhibition or other means</td> <td>*Y</td> </tr> <tr> <td>*P</td> <td>document published prior to the international filing date but later than the priority date claimed</td> <td>document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art</td> </tr> <tr> <td></td> <td></td> <td>*G</td> </tr> <tr> <td></td> <td></td> <td>document member of the same patent family</td> </tr> </table>			* Special categories of cited documents:	*T	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention	*A	document defining the general state of the art which is not considered to be of particular relevance		*E	earlier document published on or after the international filing date	*X	*L	document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone	*O	document referring to an oral disclosure, use, exhibition or other means	*Y	*P	document published prior to the international filing date but later than the priority date claimed	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art			*G			document member of the same patent family
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Date of the actual completion of the international search 14 DECEMBER 1999		Date of mailing of the international search report 25 JAN 2001																								
Name and mailing address of the ISA/US Commissioner of Patents and Trademarks Box PCT Washington, D.C. 20231 Facsimile No. (703) 305-3230		Authorized officer EDUARDO C. ROBERT Telephone No. (703) 308-0858																								

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US00/40701

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 3,757,783 A (ALLEY) 11 September 1973, see complete document.	1-9
A, P	US 6,020,380 A (KILLIAN) 01 February 2000, see complete document.	1-9